

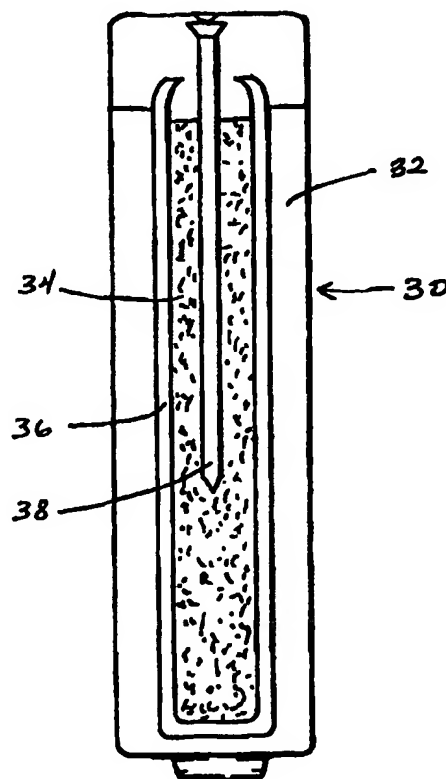


INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : H01M 4/42, 4/12	A1	(11) International Publication Number: WO 98/50969 (43) International Publication Date: 12 November 1998 (12.11.98)
(21) International Application Number: PCT/US98/08824 (22) International Filing Date: 1 May 1998 (01.05.98) (30) Priority Data: 60/045,360 2 May 1997 (02.05.97) US (71) Applicant: EVEREADY BATTERY COMPANY, INC. [US/US]; 25225 Detroit Road, Westlake, OH 44145 (US). (71)(72) Applicants and Inventors: URRY, Lewis, F. [US/US]; 36263 Butternut Ridge, Elyria, OH 44039 (US). CLASH, David, G. [US/US]; 22780 Esther Avenue, Fairview Park, OH 44126 (US). TANG, Nghia, Cong [CH/US]; 26955 Ar- bor Lane, Olmsted Township, OH 44138 (US). MANS- FIELD, Robert, N., Jr. [US/US]; 13535 South Partridge Drive, Valley View, OH 44125 (US). RANDELL, Chris [GB/GB]; 20 Woolsingham Drive, Newton Hall, Durham DH1 5SP (GB). (74) Agents: WELSH, Robert, W. et al.; Eveready Battery Com- pany, Inc., 25225 Detroit Road, Westlake, OH 44145 (US).	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	

(54) Title: ZINC SHAPES FOR ANODES OF ELECTROCHEMICAL CELLS**(57) Abstract**

An electrochemical cell (30) which contains both a cathode (32) and an anode (34). The anode contains a mixture of uniformly shaped particles (10) and electrolyte. The anode can also contain a zinc powder comprising non-uniform shaped, which can be mixed with the uniformly shaped particles of the present invention.



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

5 ZINC SHAPES FOR ANODES OF ELECTROCHEMICAL CELLS

FIELD OF THE INVENTION

10 This invention relates to the use of uniformly shaped non-mercury-added zinc particles as the anode's active material in an electrochemical cell. This invention also relates to the use of uniformly shaped and uniformly sized non-mercury-added zinc particles as the anode's active material in an electrochemical cell.

15 BACKGROUND OF THE INVENTION

Aqueous alkaline manganese dioxide zinc batteries consume zinc metal particles during electrochemical discharge. Zinc particle electrodes have the advantage of high active surface area per gram of zinc which allows high
20 average current densities without excessive polarization. The particles are made to contact a current collector. An aqueous electrolyte solution provides ionic conduction between the anode and the cathode. The aqueous electrolyte may also function as a reactant or provide reactants. The electrolyte solution is typically an aqueous alkaline electrolyte containing potassium hydroxide,
25 sodium hydroxide, or a mixture thereof.

On discharge, the zinc particles produce electricity and a reaction product consisting of zinc hydroxides, zincate ions, or zinc oxides. Generally, conventional zinc particles have had irregular shapes. The elimination of
30 mercury in the zinc anode of an electrochemical cell has resulted in poor electrical contact between the zinc particles and has led to an increase in corrosion. One method used to overcome poor electrical contact has been to use higher concentration of zinc in the anode. However, this reduces the space

5 available for reaction products. Another method used to overcome poor electrical contact has involved concentrating the zinc by creating connected agglomerates of zinc. Patents and applications that describe the use of agglomerates include European Patent Application EP 414,990 A1; U.S. Patent No. 4,963,447 to Nishimura et. al.; and U.S. Patent No. 3,884,722 to
10 Tucholski. Unfortunately, using the gelling agents described in these patents can impair the cell's discharge characteristics by inhibiting the reaction efficiency of the zinc.

Others in attempts to reduce anode gassing have compressed the zinc
15 particles such as in Vu et al. ; U.S. Patent 4,743,185. Others have used zinc powders of a generally spherical shape such as in Japanese unexamined patent publication 4-296451. However, this publication teaches zinc apparent densities greater than 3.5 g/cc with anode zinc concentrations as high as 75 weight percent.

20

In order to achieve the highest electrochemical activity from the zinc particles with the highest particle-to-particle contact and electroconductivity, it is desirable to have an anode comprising zinc particles having uniform morphology and/or size. It would also be desirable to have zinc particles
25 having the same porosity, chemical composition, reactivity, morphological, topological, and physical properties. Battery systems in which zinc particles of this invention may be used include alkaline manganese dioxide-zinc cells, zinc-air button cells, and the like, including primary and secondary batteries.

5

SUMMARY OF THE INVENTION

This invention is an electrochemical cell comprising a cathode, an aqueous alkaline electrolyte, and a non-mercury-added anode comprising
10 uniformly shaped zinc particles.

In another aspect, this invention is an electrochemical cell comprising a cathode, an aqueous alkaline electrolyte, and a non-mercury-added anode comprising uniformly shaped zinc particles having uniform size.

15

The electrochemical cell of the present invention includes the uniformly shaped zinc particles having a tap density less than 2.5 g/cc, preferably having a tap density less than 2.0 g/cc, very preferably having a tap density less than 1.5 g/cc, and most preferably having a tap density less than
20 1.3 g/cc.

In the electrochemical cell of the present invention the uniformly shaped zinc particles comprise fifty percent or less of the zinc in the anode, preferably the uniformly shaped zinc particles comprise twenty percent or less
25 of the zinc in the anode, very preferably the uniformly shaped zinc particles comprise fifteen percent or less of the zinc in the anode, and most preferably the uniformly shaped zinc particles comprise ten percent or less of the zinc in the anode.

30 In the electrochemical cell of the present invention containing the uniformly shaped zinc particles the zinc weight concentration of the anode is less than 70 percent, preferably the zinc weight concentration of the anode is less than 68 percent, very preferably the zinc weight concentration of the

5 anode is less than 66 percent, most preferably the zinc weight concentration of the anode is less than 64 percent or below 62 percent.

In a preferred embodiment of the electrochemical cell of the present invention the uniformly shaped particles have the length, width and thickness
10 dimensions in the range of 0.06, 0.06 and 0.0025 to 0.01, 0.01 and 0.0004 inches, most preferably the length, width and thickness dimensions are in the range of 0.04, 0.025 and 0.0008 inches to 0.02, 0.025, and 0.0008 inches.

In a preferred embodiment of the invention the zinc weight
15 concentration of the anode is less than 67 percent, the uniformly shaped zinc particles comprise twenty percent or less of the zinc in the anode, and the uniformly shaped zinc particles having a tap density less than 2.5 g/cc.

DETAILED DESCRIPTION OF THE INVENTION

20

Generally, the present invention is an electrochemical cell comprising the novel use of a non-mercury-added anode comprising uniformly shaped zinc particles. According to this invention, the use of zinc particles having a uniform morphology increases the performance of the zinc anode in an
25 electrochemical cell.

The inventors have found that the use of uniformly shaped zinc particles can have one or more of the following advantages. First, the surface area/volume ratio can be increased. Second, the current carrying matrix of zinc
30 particles is easier to maintain. Third, the zinc particles' high current discharge efficiency can be improved. Fourth, the anode's sensitivity to shock and vibration can be reduced or eliminated. Fifth, the quantity of zinc packing needed can be minimized. Sixth, conductivity within the anode is improved.

5 Seventh, processability, flowability, and dispensability of the anode is improved. Eighth, the use of large particle gelling agents can be eliminated. Ninth, the quantity of gelling agent used in the anode can be reduced. Tenth, the zinc-to-electrolyte ratio is improved. Eleventh, zinc-to-zinc contact is improved. Twelfth, the distribution of zinc in the gelled electrolyte is
10 improved.

The expression "uniform shape" or "uniformly shaped" is intended to mean individual particles having substantially consistent morphology. This definition is contrasted against the prior art zinc powders with consistently
15 irregularly shapes, few of which are similar to each other. To have a uniform shape as described and claimed, each of the particles shape factor must be substantially similar to all other particles. Therefore for example, if the uniform shape is flakes, then all of the particles must be flakes. In order to produce uniformly shaped particles, control of the forming or classification
20 process should be used to ensure this required consistent particle shape.

The shapes of zinc particles can be determined by examining the particles with a scanning electron microscope. Preferably, the zinc particles of this invention will also have substantially consistent surface finishes and
25 porosity characteristics.

As found in this invention, the shape of zinc particles is critical to electrochemical activity and anode processing. The size can also be critical to electrochemical activity and anode processing. When the particles are
30 irregularly shaped or too large, processability of the anode is difficult. However, when the particles are too small, there can be increased ionic and electrical resistance.

5 More preferably, the zinc particles of this invention will have a uniform morphology and a uniform size. The use of uniformly shaped zinc particles with uniform size has the advantages of providing uniform bulk density and consistent particle-to-particle contact. The shape and size of the zinc particle can be optimized depending on the application. For uniformly
10 shaped spherical zinc particles provide enhanced processability of the anode. Zinc particles having a "dog-bone" morphology provide an advantage when used in continuous drain applications. However, as the particle size gets larger, flowability of the anode may decrease. Also, although the large particles provide relatively larger areas of contact, the smaller particles are preferred
15 due to enhanced zinc-to-zinc contact and improved flowability.

 The zinc particles of this invention can also have substantially similar porosity characteristics. By "porosity" is meant a particle having intraparticle void volume greater than zero. A porous zinc particle can also comprise
20 interconnected pores. The total porosity of the zinc is preferably greater than 25 percent porosity and, more preferably, greater than 50 percent porosity as determined by bulk or tap density analyses. An advantage to having a highly porous zinc electrode is the ability to store electrolyte in the pores of the electrode and then use this electrolyte to replenish the water and hydroxyl ions
25 which are consumed during discharge of the cell. Porous zinc particles provide a high surface area-to-volume ratio which is important in high drain rate applications. Particle morphologies having intraparticle void volume can be shaped as a sponge, honeycomb, foam, tube, and the like.

30 The zinc particles of this invention can also be solid zinc particles with no internal void volume. Particle morphologies having no intraparticle void volume can be shaped as a flake, snake, filament, donut, sphere, dogbone, biscuit, or dendrite. These particles may be corrugated. Suitable morphologies

5 include tentacle-like, helix, or faceted. Flakes can have the shapes of discs, ribbons, clamshells, triangles, squares, troughs, rhomboids, rectangles, boomerangs, and the like. "Snake" morphology can have the shapes of rods, cylinders, carrots, needles, worms, or bananas. In general, "snake" morphology would include those shapes having an approximately round cross-section, not
10 necessarily uniform along its length, where the ratio of length to average diameter is greater than 1 and wherein the axis is defined as a line connecting a locus of points through the centers of all cross-sections. This axis of round cross-section is not necessarily straight. Filament morphology resembles wire, hair, thread, spaghetti, fibrils, fiber, ribbon, or turnings. The zinc particles can
15 also have "donut" or torus morphology. The zinc particles can also have spheroid morphology, such as spheres, tadpoles, teardrops, pears, or squash-like shapes. Zinc particles having "dog bone" morphology can best be described as those particles having an approximately round cross-section, not necessarily uniform along its length, where the ratio of length to average
20 diameter is greater than 1, the axis is not necessarily straight, and the diameters of both ends are greater than the average diameter along the axis. A biscuit morphology is described as a briquette, flattened sphere, or cookie-like morphology. A dendrite morphology resembles feathers or snowflakes. A corrugated morphology resembles waffles, or mesh screens. A tentacle
25 morphology resembles plant roots, octopus, or squid. A helix morphology resembles springs, scrolls, or turnings. A faceted morphology resembles pyramid or multifaceted objects.

Surface finishes that are suitable for use with particles of this invention
30 include: smooth, mossy, porous, grooved, dendritic and waffle-like.

The uniformly shaped zinc particles of this invention can be formed in a variety of ways. For example, U.S. Patent No. 4,154,284 describes a process

5 of producing flake particles having a small length-to-width or small
length-to-thickness ratio by extraction from a molten pool or unconfined drop.
The process utilizes a rotatable heat-extracting member having a serrated
circular peripheral edge with each serration having a leading surface angularly
shaped to contact the surface of the molten material. This process is capable of
10 controlling the shape, size, and thickness of the final flake product.
Furthermore, the flakes will be "substantially uniformly shaped." The precise
shape and dimensions of the individual flake particles can be controlled by
changing one or more of the following process parameters: the angle at which
the serrated edge contacts the molten material; the height of the serration, the
15 length of the serration, the diameter of the rotating member; the rate at which
the molten flake is cooled; the atmosphere (air vs. vacuum); etc.

A process suitable for making particles having "snake" like
morphology (as described above) is disclosed in U. S. Patent No. 3,871,439.
20 This patent discloses a method of making fine filamentary material directly
from molten material by rotating the peripheral edge of a machine threaded
cylindrical heat-exchanging member in contact with the surface of the molten
material. Small fibers (i.e., less than 100 microns in length) are readily
produced by this process. The length of the fibers can be controlled by
25 incorporating indentations on the extremities of the projections that define the
surface of the cylindrical heat-exchanging member.

A process for manufacturing "uniform particulate" is disclosed in U. S.
Patent No. 4,385,013. This process produces particulate directly from a supply
30 of molten material by the use of a rotating member having discrete serrations
in the periphery thereof. The leading surfaces of the serration contact the
molten material and propel a portion into a cavity formed by the surfaces of
the serration while under the effect of a surface of a dam means, in proximity

5 to which the serrations pass. The dam means is immersed in the supply of molten material adjacent to the periphery of the rotating member. Provided that each of the serrations on the rotating member is identical, the particulate produced will be uniform in shape and size. By changing the shape of the serrations, the shape of the particulate can be altered to any one of a wide
10 variety of shapes. By way of example, three of the shapes disclosed include: quadrahedron, a five-sided triangular particle, and a six-sided particle.

The zinc particles of this invention can be purchased from known zinc manufacturers such as Transmet Corporation of Columbus, Ohio, U.S.A. or
15 Eckart-PM-Laboratory of Vetroz, Switzerland.

EXAMPLES

Two different zinc shapes were added to a conventional zinc powder
20 composition in different amount using a total zinc weight percent of the anode in the range of 62-64%. A control was used with 100 wt. % conventional zinc powder and a total zinc weight percent of the anode in the range of 69-71%.

Each of the experimental cells were manufactured so that the cathode
25 theoretical Ampere hour value and anode theoretical Ampere hour value were kept constant regardless of the total zinc content. This was accomplished by using a larger cathode ID for the 62-64 weight percent total content anodes.

The experiments were done in three parts:

30 1. Shock testing - The short circuit amperage (SCA) of the cell was taken before and after drop testing (procedure defined below).

2. Performance testing - 1A continuous, 1A Photoflash, and 3R9 Continuous.

- 5 3. Impedance testing - the impedance of the anodes were measured on an Autolab PGSTAT20 using FRA software from EchChemie.

Example 1:

- 10 The tap density of all zinc samples were first measured according to the following procedure:

1. Zinc is put in a pre-weighed 25ml measuring cylinder.
2. The zinc is then tapped on a rubber bung whereby the level of the
15 zinc falls.
3. Extra zinc is added in stages until the zinc level is exactly 25ml in the measuring cylinder.
4. The measuring cylinder is weighed again.
5. The weight of zinc which occupies 25ml can then be calculated.
- 20 6. The Tap density is the weight of zinc which occupies 25ml / 25ml

The results were calculated and are tabulated in Table 1.

Table 1

Zinc type	Tap Density (g/cc)
Needles	3.6
Conventional powder	3.4
Flakes	1.3

25

Example 2:

Numerous groups of cells were manufactured with two different cell formulations.

5 Formulation 1

Cathode Ah = 2.7

EMD : C = 12:1

ID (mm) = 8.73

Anode Ah = 3.3

10 Zinc Content = 69%

H₂O/EMD = 0.24Formulation 2

Cathode Ah = 2.7

15 EMD : C = 17:1

ID (mm) = 9.00

Anode Ah = 3.3

Zinc Content = 63

H₂O/EMD = 0.24

20

Common Features

EMD GHU by TOSOH

Graphite Superior Graphite

KOH Conc. 35%

25 Can Coating LB1099 by Timcal

	<u>Zinc Types</u>	<u>Alloys</u>	<u>Designation</u>	<u>Supplier</u>
	Zinc Powder	BIC	Powder	Big River Zinc
	Zinc Flake	BIC	K308	Transmet of Ohio
30	Zinc Flake	NOT BIC	K109	Transmet of Ohio
	Zinc Needles	BIC	Doralloy 107 GN	Eckart-Doral of Switzerland

5 Example 3:

The cells were tested according to the following drop test method. The results of the testing are compiled in Table 2, and presented graphically in Figure 7.

- 10 1. The SCA (Flash Amps) of the cell is measured 1 week after manufacture.
2. The cell is allowed to fall down a vertical shaft. The length of the shaft is 1m. To start the drop the cell is rolled so that it always falls on its side rather than the ends.
3. The SCA (Flash Amps) of the cell is measured immediately after being
- 15 rolled down the vertical shaft.

Table 2

Anode comp.	Amperage before drop	Amperage after drop	Amperage maintenance (%)
100/0/70	16	4	25
100/0/63	7	0	0
95/5/63	18	5	28
90/10/63	19	9	47
85/15/63	21	9	43

- 20 Table 2 shows the results of the drop test for various anode concentrations. The concentrations should be interpreted as (% of zinc powder/ % of zinc flake/ anode zinc weight percent). Amperage maintenance is amperage after drop divided by amperage before drop.

- Referring now to Figure 7, there is shown a graphical representation of
- 25 the results presented in Table 2. Both Table 2 and this figure show the superior

- 5 amperage maintenance results achieved in low zinc anode concentrations by the addition of uniform shaped zinc to the anode.

Example 4:

- 10 The cells formed in example 2 were tested according to standard industry duration tests. The results of the tests are presented both in Table 3 and in Figure 8. The results clearly show the advantages supplied to batteries using the present invention.

15

Table 3

	Relative to 100/0/70			Test Data		
	3.9 ohm 1h/d 0.8V	10 ohm 1h/d 0.9V	43 ohm 1h/d 0.9V	3.9 ohm 1h/d 0.8V	10 ohm 1h/d 0.9V	43 ohm 1h/d 0.9V
100/0/70	100%	100%	100%	6.5	16.9	77
100/0/62	111%	96%	109%	7.2	16.2	84
95/05/62	111%	101%	112%	7.2	17.1	86
90/10/62	114%	103%	118%	7.4	17.4	91
85/15/62	118%	104%	116%	7.7	17.6	89
80/20/62	120%	105%	112%	7.8	17.8	86

Example 5:

- In this example cells constructed in example 2 were subjected to impedance testing. This method of testing is known in the art and was
 20 conducted using an Autolab PGSTAT20 using FRA software from EchChemie. The results are shown in Table 4 below.

5

Table 4

Total Zinc Content	KOH	% Flake	10KHz		1KHz		100Hz		10Hz		1Hz		0.1Hz		0.01Hz	
			z'	z''	z'	z''	z'	z''	z'	z''	z'	z''	z'	z''	z'	z''
70.5%	35%	0%	0.3	-0.1	0.3	0.1	0.4	0.8	3.2	5.8	15	5	18	2	19	2
63.5%	35%	0%	0.8	0.5	1.7	2.5	3.9	28	105	173	391	132	487	65	501	50
63.5%	35%	5%	0.3	0	0.3	0.1	0.4	0.7	2.2	5.6	17	8	22	3	22	2
63.5%	35%	10%	0.2	-0.1	0.3	0	0.3	0.4	1.7	3	8	3	10	1	11	1
63.5%	35%	15%	0.2	-0.1	0.2	0	0.3	0.4	1.4	2.9	9	3	10	1	11	1

In the Table, z' is the real impedance, and z'' is the imaginary impedance.

10

The results show that as the total zinc content of the anode is decreased the impedance dramatically increases. This is indicative that the cell would be inoperative when the total zinc content is dropped much below 70 weight percent.

15

However when the uniform shapes are added to the anode the total zinc concentration of the anode can be decreased. An optimal level of flake appears at 10%.

20

It is well known that if the total zinc content of a 0Hg zinc anode falls below 67% the resistance to shock deteriorates. This has been demonstrated in example 3. After 1m drop testing 62-64% total zinc content paste containing ONLY normal zinc shows 0A SCA. It is possible to recover the shock resistance of 62-64% total zinc content paste by adding zinc flake. However the addition of zinc needles does not produce desirable results.

25

5 For flakes, as the flake content is increased the usability of the anode gel decreases. The best compromise between shock resistance and usability is ~10% zinc flake content. However this optimal point is dependant upon the type of uniformly shaped particle and will differ with different geometries.

10 At 10% flake content for 62-64% total zinc content gel performance reaches an overall maximum for all performance tests (from very high rate to very low rate). A typical very high rate test is 1A / Cont. / 1V0. A typical very low rate test is 43R0 / 4h / 0V9. However the incorporation of Dorally 07CN needles though has no beneficial effect on 62-64% total zinc content
15 pastes.

 The main difference between the flakes and the needles is the tap density. The tap density of the zincs used in this study are given above in Table 1. The flake has a very low tap density. And it is believed that this is the
20 key to why the flakes work with 62-64% total zinc content gel but the needles do not. The low density flakes make a matrix through the anode gel which gives resistance to shock. Also very high rate 1A performance is boosted. The needles are too high a density to make the required matrix, so shock resistance & 1A performance suffer.

25

 So the advantage of low tap density zinc is that it allows a matrix to be formed in low total zinc content anode gel. This matrix gives good shock resistance.

30

 Adding low tap density zinc allows low total zinc content to be used in alkaline manganese. As the Hg level was reduced in alkaline manganese zinc anodes the total zinc content of the anode paste has been raised. Typically in the days of 6% Hg the total zinc content of AA anode paste was as low as

5 64.75%. With cells used today with conventional 0% Hg zinc powders, the
typical total zinc content is 70%. Attempts of local densification have been
made by adding nuggets. The use of low tap density zinc allows total zinc
contents of 62-64% to be used with 0% Hg. The extra electrolyte bound in
such a gel leads to improved efficiency. This in turn leads to improved
10 performance.

DESCRIPTION OF THE DRAWINGS

15

Figure 1 is a scanning electron micrograph of zinc particles having a uniform biscuit shape of the present invention.

Figure 2 is a scanning electron micrograph of zinc particles having a uniform needle shape of the present invention.

20 Figure 3 is a scanning electron micrograph of zinc particles having a uniform, ribbon shape of the present invention.

Figure 4 is a representative drawing of a scanning electron micrograph of zinc particles having a uniform, flake shape of the present invention.

25 Figure 5 is a representative drawing of a scanning electron micrograph of zinc particles having a non-uniform shape. This type of zinc particle is typical of the prior art and is normally referred to as zinc powder.

Figure 6 is a cross-sectional view of an electrochemical cell of the present invention.

30 Figure 7 is a graph showing the results of a drop test described in the examples specifically in relation with Table 2.

Figure 8 is a graph showing the results of cell testing as described in the examples.

5

DETAILED DESCRIPTION OF THE DRAWINGS

Referring now to the drawings and specifically to Figure 4, there is show a uniformly shaped zinc particle 10 of the present invention. Each of the
10 other particles are substantially the same size and geometry as the particle 10. Contrast this with the prior art zinc particle 20 shown in Figure 5, which is dissimilar relative to the other particles in the figure.

Referring now to Figure 6, there is shown an electrochemical cell 30 of the present invention. The cell 30 contains both a cathode 32 and an anode 34.
15 The anode contains a mixture of uniformly shaped particles and electrolyte. The electrolyte is preferably potassium hydroxide, however other alkaline or nonalkaline electrolyte are suitable alternatives. The anode can also contain a gelling substance, several of which are well known in the art. The anode can also contain a zinc powder comprising non-uniform shaped, which can be
20 mixed with the uniformly shaped particles of the present invention.

The cell 30 also includes a separator 36 which electrically insulates the anode from the cathode, while allowing for ionic diffusion there through. The cell 30 also contains a current collector 38 which acts to collect the electrons
25 from the anode 344 and transport them to a external apparatus so that work can be performed.

- 5 Changes in the composition of the anode materials, the processes, the structure, and components of the cells may be made without departing from the scope of the present invention as defined in the following claims.

CLAIMS

10

What is claimed is:

1. An electrochemical cell comprising a non-mercury-added anode comprising uniformly shaped zinc particles.
2. An electrochemical cell of claim 1 wherein the uniformly
15 shaped zinc particles have a tap density less than 2.5 g/cc.
3. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles have a tap density less than 2.0 g/cc.
4. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles have a tap density less than 1.5 g/cc.
- 20 5. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles have a tap density less than 1.3 g/cc.
6. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles comprise fifty percent or less of the zinc in the anode.
- 25 7. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles comprise fifty percent or less of the zinc in the anode.
8. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles comprise twenty percent or less of the zinc in the
30 anode.
9. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles comprise fifteen percent or less of the zinc in the anode.

- 5 10. An electrochemical cell of claim 1 wherein the uniformly shaped zinc particles comprise ten percent or less of the zinc in the anode.
11. An electrochemical cell of claim 1 wherein the zinc weight concentration of the anode is less than 70 percent.
- 10 12. An electrochemical cell of claim 1 wherein the zinc weight concentration of the anode is less than 68 percent.
13. An electrochemical cell of claim 1 wherein the zinc weight concentration of the anode is less than 66 percent.
- 15 14. An electrochemical cell of claim 1 wherein the zinc weight concentration of the anode is less than 64 percent.
15. An electrochemical cell of claim 1 wherein the zinc weight concentration of the anode is less than 62 percent.
16. An electrochemical cell of claim 1 wherein the uniformly shaped particles have the length, width and thickness dimensions of 0.06, 0.06 and 0.0025 inches or less.
- 20 17. An electrochemical cell of claim 2 wherein the zinc weight concentration of the anode is less than 67 percent.
18. An electrochemical cell of claim 2 wherein the uniformly shaped zinc particles comprise twenty percent or less of the zinc in the anode.
- 25 19. An electrochemical cell of claim 17 wherein the uniformly shaped zinc particles comprise twenty percent or less of the zinc in the anode

FIGURE 1

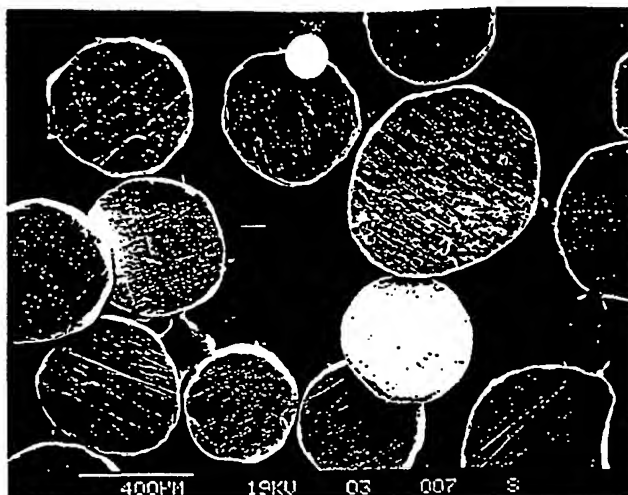


FIGURE 2

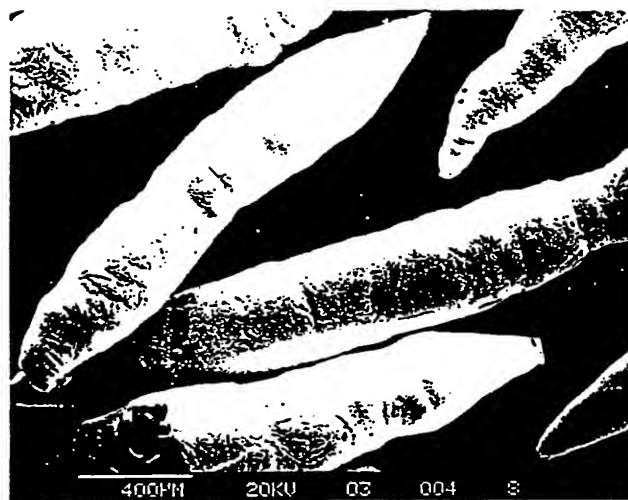
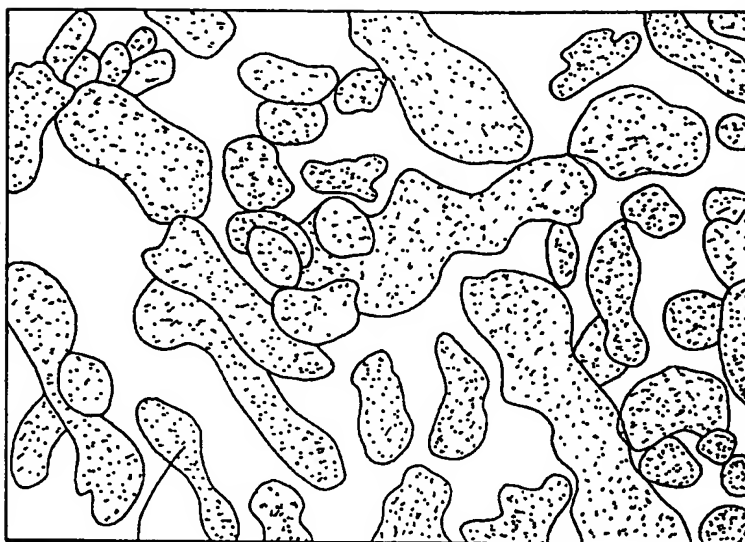


FIGURE 3



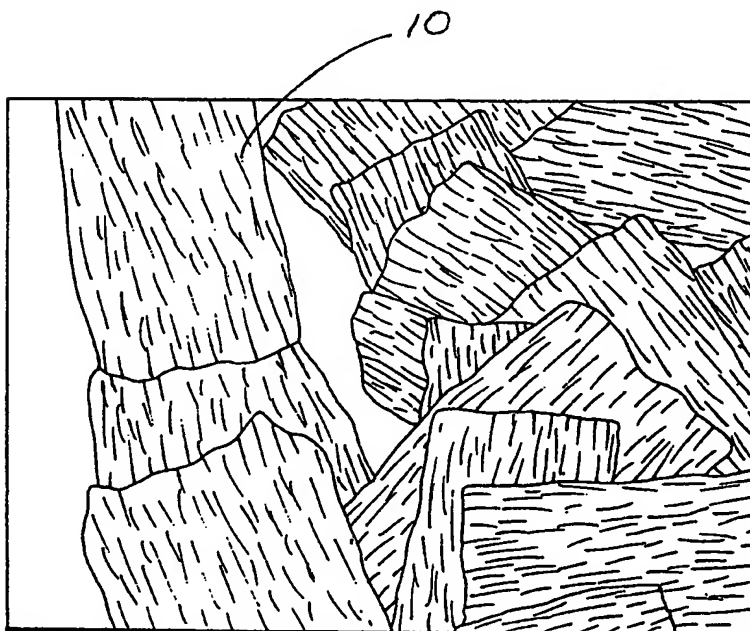
2/5

Fig. 5



20 ZINC POWDER

Fig. 4



10 ZINC FLAKES

3/5

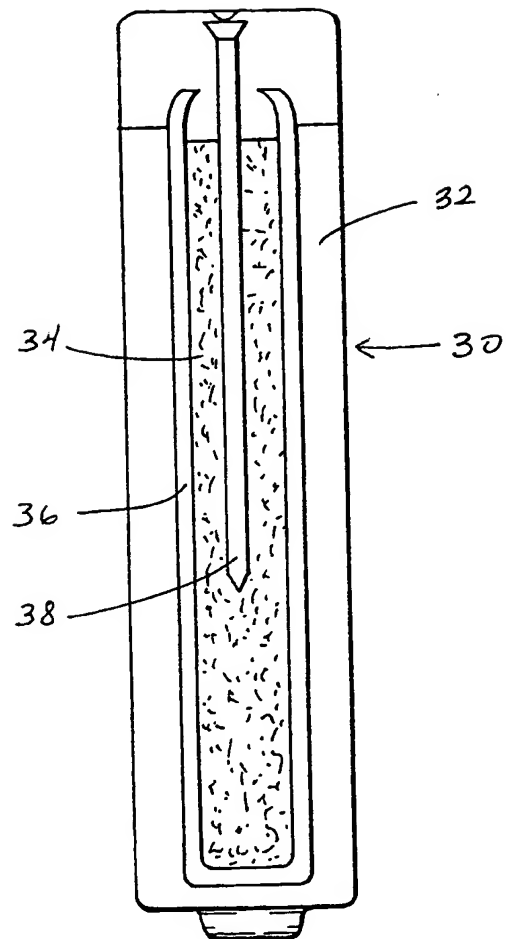
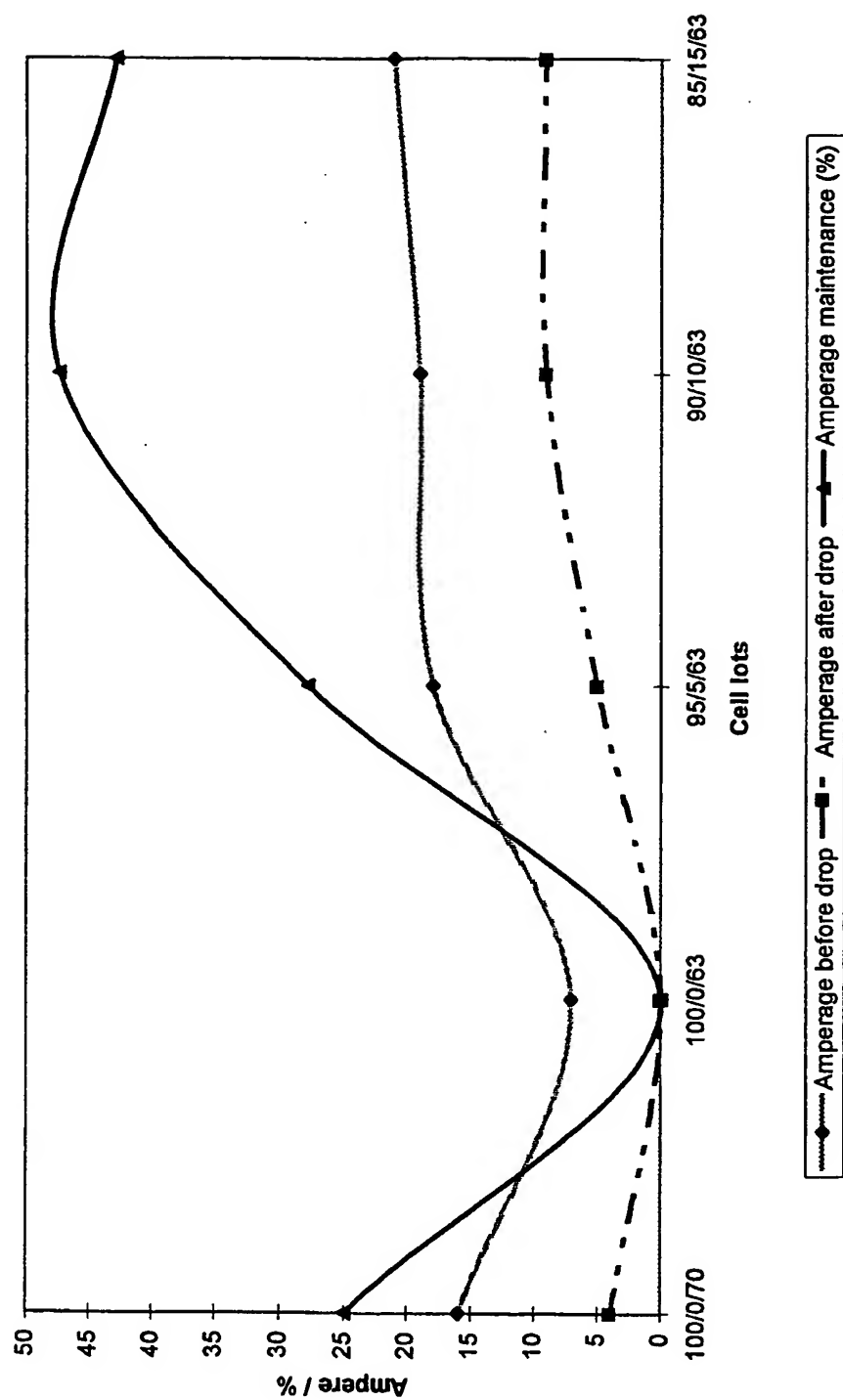


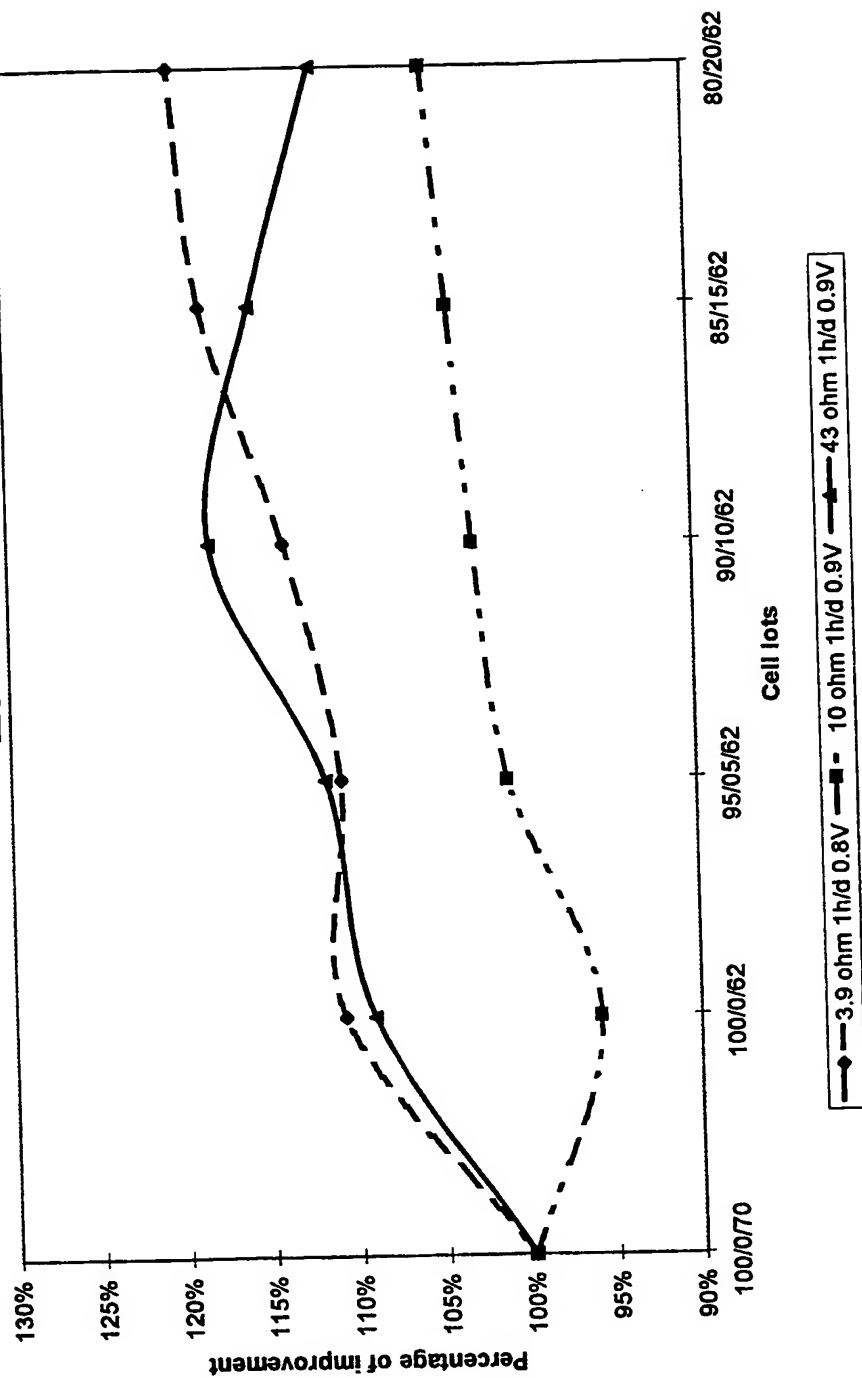
Fig. 6.

4/5

Figure 7

5/5

Figure 8



INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/08824

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 H01M4/42 H01M4/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 H01M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	WO 98 20569 A (EVEREADY BATTERY INC) 14 May 1998 see page 5, line 17 - line 19; claims 1,10,18; table 1 ---	1-19
X	US 4 606 869 A (SHOWAK WALTER) 19 August 1986 see column 2, line 13 - line 18; table 1 ---	1
X	PATENT ABSTRACTS OF JAPAN vol. 008, no. 070 (E-235), 3 April 1984 & JP 58 218760 A (TOSHIBA DENCHI KK), 20 December 1983 see abstract --- -/--	1

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

5 August 1998

Date of mailing of the international search report

13/08/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Andrews, M

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 98/08824

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>PATENT ABSTRACTS OF JAPAN vol. 096, no. 002, 29 February 1996 & JP 07 254406 A (TOSHIBA BATTERY CO LTD), 3 October 1995 see abstract</p> <p>-----</p>	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/08824

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9820569	A	14-05-1998	NONE	
US 4606869	A	19-08-1986	NONE	

THIS PAGE BLANK (USPTO)